Attorney Docket No. 22112(3)

CLAIMS

What is claimed:

A stable formulation of a biologically active protein useful for aerosol delivery to the respiratory tract of a patient in need of treatment comprising:

- (a) a carrier liquid comprising from about 10% to from about 100% V/V water and from about 0% to from about 90% V/V of an organic liquid;
- (b) a biologically effective amount of a protein suspended or dissolved in a carrier liquid; and
- (c) a stabilizing effective amount of a derivatized carbohydrate stabilizing agent suspended or dissolved in said carrier liquid.
- 2. A stable suspension according to claim 1 wherein said formulation contains from about 0.1% to about 5.0% W/V of a pharmaceutically acceptable excipient.
- 3. A stable formulation according to claim 1 wherein said biologically active protein is selected from the group comprising enzymes, antibodies, antigens, hormones and cytokines.
- 4. A stable formulation according to claim 3 wherein said therapeutically active protein is a hormone.
 - 5. A stable formulation according to claim 4 wherein said therapeutically active protein is insulin.
- 6. A stable formulation according to claim 3 wherein said therapeutically active protein is a cytokine.
- 7. A stable formulation according to claim 6 wherein said therapeutically active protein is Factor VIII.

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- 8. A stable formulation according to claim 1 wherein said carrier liquid contains from about 20% to from about 100% water V/V.
- 9. A stable formulation according to claim 8 wherein said carrier liquid comprises about 50% water and about 50% organic solvent.
 - 10. A stable formulation according to claim 1 wherein said organic liquid is ethanol, isopropyl alcohol, butanol, isobutanol, perfluorocarbons, glycerol, polyethylene glycol, propylene glycol, or combinations thereof.
 - 11. A stable formulation according to claim 10 wherein said organic liquid is ethanol, glycerol, polyethylene glycol, propylene glycol, or combinations thereof.
 - 12. A stable formulation according to claim 1 wherein said sugar moiety is selected from the group consisting of trehalose, sucrose, glucose, maltose, and galactose.
 - 13. A stable formulation according to claim 1 wherein said protein is suspended in the carrier liquid.
 - 14. A stable formulation according to claim 13 wherein the particle size of said protein in suspension is from about 0.01 μ to about 10.0 μ .
 - 15. A stable formulation according to claim 14 wherein the particle size of said protein in suspension is from about 5.0 μ to about 10.0 μ.
 - 16. A stable formulation according to claim 15 wherein the particle size of said protein in suspension is from about 0.01 μ to about 3.0 μ .
 - 17. A stable formulation according to claim 2 wherein said formulation contains from about 0.1% to about 5.0% of a pharmaceutically acceptable excipient.

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- 18. A stable formulation according to claim 1 wherein said protein is dissolved in the carrier liquid.
- 19. A stable formulation according to claim 18 wherein said formulation contains from about 0.1% to about 5.0% of a pharmaceutically acceptable excipient.
 - 20. A stable formulation of a biologically active protein useful for aerosol delivery to the respiratory tract of a patient in need of treatment comprising:
 - (a) a carrier liquid which is from about 20% to from about 30% V/V water and from about 70% to from about 80% V/V of ethanol;
 - (b) a biologically effective amount of a protein suspended or dissolved in a carrier liquid; and
 - (c) a stabilizing effective amount of a derivatized carbohydrate stabilizing agent elected from the group consisting of C8-trehalose, C16-trehalose, C8-glycopyranoside and C12-glucopyranoside.
 - 21. A stable formulation of a biologically active protein useful for aerosol delivery to the respiratory tract of a patient in need of treatment comprising:
 - (a) a carrier liquid which is an aqueous liquid;)
 - (b) a biologically effective amount of a protein suspended or dissolved in said carrier liquid; and
 - (c) a stabilizing effective amount of a derivatized carbohydrate stabilizing agent elected from the group consisting of C8-trehalose, C16-trehalose, C8-glycopyranoside and C12-glucopyranoside.